Title: Deglycosylated Kringle 1-3 Region Fragments of Plasminogen and Methods of Use

Amendment and Response to Office Action

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AMENDMENTS TO THE CLAIMS

This listing of the claims will replace all prior versions, and listings, of claims in the application:

- (Currently Amended) A composition comprising a pharmaceutically 1. acceptable carrier and a protein consisting of a deglycosylated kringle 1-3 region fragment of a plasminogen protein, wherein the deglycosylated kringle 1-3 region fragment lacks one or more two carbohydrate mojeties found in mojeties linked to naturally glycosylated forms of the fragment, wherein the deglycosylated kringle 1-3 region fragment has antiangiogenic activity, and wherein the deglycosylated kringle 1-3 region fragment and a glycosylated form of the fragment are at a ratio of 100:0.
- 2. (Previously Presented) The composition of claim 1, wherein the deglycosylated kringle 1-3 region fragment lacks a bisialylated-biantennary glycan.
- (Previously Presented) The composition of claim 1, wherein the 3. deglycosylated kringle 1-3 region fragment lacks an N-linked carbohydrate moiety.
- (Previously Presented) The composition of claim 1, wherein the 4. deglycosylated kringle 1-3 region fragment lacks a carbohydrate chain at an amino acid position corresponding to an N-glycosylation site of human plasminogen.

5. (Cancelled)

- (Previously Presented) The composition of claim 1, wherein the deglycosylated kringle 1-3 region fragment begins at approximately amino acid 87 of human plasminogen.
- (Previously Presented) The composition of claim 1, wherein the deglycosylated kringle 1-3 region fragment amino acid sequence is shown in SEQ ID NO:2.

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(Previously Presented) The composition of claim 1, wherein the deglycosylated kringle 1-3 region fragment is produced recombinantly.

6 9. (Previously Presented) The composition of claim 1, wherein the deglycosylated kringle 1-3 region fragment has an amino acid substitution at amino acid position corresponding to the N-glycosylation site of human plasminogen.

10-14. (Cancelled)

15. (Previously Presented) The composition of claim 1, wherein the deglycosylated kringle 1-3 region fragment has antiangiogenic activity in vitro.

(Previously Presented) The composition of claim 1, wherein the deglycosylated kringle 1-3 region fragment has antiangiogenic activity in vivo.

17-26 (Cancelled)

27. (Previously Presented) A deglycosylated kringle 1-3 region fragment of a plasminogen protein, wherein the deglycosylated kringle 1-3 region fragment amino acid sequence is shown in SEQ ID NO:2.

28. (Cancelled)

29: (Previously Presented) The composition of claim 40, wherein the amount of the naturally glycosylated kringle 1-3 region fragment present in the composition is smaller than the amount of the deglycosylated kringle 1-3 region fragment present in the composition.

30-34. (Cancelled)

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35. (Previously Presented) The composition of claim 39, wherein the deglycosylated kringle 1-3 region fragment is produced recombinantly.

36. (Cancelled)

37. (Previously Presented) The composition of claim 39, wherein the deglycosylated kringle 1-3 region fragment has antiangiogenic activity in vitro.

38. (Previously Presented) The composition of claim 39, wherein the deglycosylated kringle 1-3 region fragment has antiangiogenic activity in vivo.

39. (Previously Presented) A composition comprising a pharmaceutically acceptable carrier and a protein consisting of a deglycosylated kringle 1-3 region fragment of a plasminogen protein wherein the deglycosylated kringle 1-3 region fragment lacks one or more carbohydrate moieties linked to naturally glycosylated forms of the fragment, wherein the deglycosylated kringle 1-3 region fragment has antiangiogenic activity, and wherein the deglycosylated kringle 1-3 region fragment amino acid sequence is shown in SEQ ID NO:2.

(Previously Presented) The composition of claim 39, further comprising a protein consisting of a naturally glycosylated kringle 1-3 region fragment of a plasminogen protein.

(Currently Amended) The <u>deglycosylated kringle 1-3 region fragment</u> composition of claim 27, wherein the deglycosylated kringle 1-3 region fragment is produced recombinantly.

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(Currently Amended) The <u>deglycosylated kringle 1-3 region fragment</u> ecomposition of claim 27, wherein the deglycosylated kringle 1-3 region fragment has antiangiogenic activity in vitro.

(Currently Amended) The <u>deglycosylated kringle 1-3 region fragment</u> emposition of claim 27, wherein the deglycosylated kringle 1-3 region fragment has antiangiogenic activity in vivo.